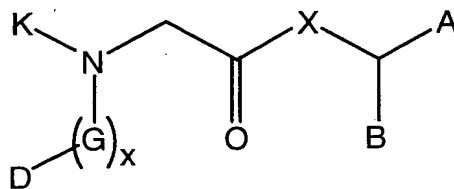


CLAIMS

1. A compound having formula (I):



(I)

and pharmaceutically acceptable derivatives thereof,  
wherein:

X is O, S, C(R<sup>1</sup>)<sub>2</sub> or NR<sup>1</sup>;

10 A, B and R<sup>1</sup> are independently E, (C<sub>1</sub>-C<sub>10</sub>)-straight or branched alkyl, (C<sub>2</sub>-C<sub>10</sub>)-straight or branched alkenyl or alkynyl, or (C<sub>5</sub>-C<sub>7</sub>)-cycloalkyl or cycloalkenyl; wherein 1 or 2 hydrogen atoms in said alkyl, alkenyl or alkynyl are optionally and independently replaced with E, (C<sub>5</sub>-C<sub>7</sub>)-  
15 cycloalkyl or cycloalkenyl; and wherein 1 to 2 methylene (-CH<sub>2</sub>-) groups in said alkyl, alkenyl, or alkynyl groups are optionally and independently replaced by -O-, -S-, -S(O)-, -S(O)<sub>2</sub>-, =N-, -N= or -N(R<sup>3</sup>)-;

or B and R<sup>1</sup> are independently hydrogen;

20 wherein R<sup>3</sup> is selected from hydrogen, (C<sub>1</sub>-C<sub>4</sub>)-straight or branched alkyl, (C<sub>3</sub>-C<sub>4</sub>)-straight or branched alkenyl or alkynyl, or (C<sub>1</sub>-C<sub>4</sub>) bridging alkyl, wherein said bridge is formed between the nitrogen atom to which said R<sup>3</sup> is bound and any carbon atom of said alkyl,  
25 alkenyl or alkynyl to form a ring, and wherein said ring is optionally benzofused;

wherein E is a saturated, partially saturated or unsaturated, or aromatic monocyclic or bicyclic ring system, wherein each ring comprises 5 to 7 ring atoms

independently selected from C, N, O or S; and wherein no more than 4 ring atoms are selected from N, O or S;

wherein 1 to 4 hydrogen atoms in E are optionally and independently replaced with halogen,  
5 hydroxyl, hydroxymethyl, nitro, SO<sub>3</sub>H, trifluoromethyl, trifluoromethoxy, (C<sub>1</sub>-C<sub>6</sub>)-straight or branched alkyl, (C<sub>2</sub>-C<sub>6</sub>)-straight or branched alkenyl, O-[(C<sub>1</sub>-C<sub>6</sub>)-straight or branched alkyl], O-[(C<sub>3</sub>-C<sub>6</sub>)-straight or branched alkenyl], (CH<sub>2</sub>)<sub>n</sub>-N(R<sup>4</sup>)(R<sup>5</sup>), (CH<sub>2</sub>)<sub>n</sub>-NH(R<sup>4</sup>)-(CH<sub>2</sub>)<sub>n</sub>-Z,  
10 (CH<sub>2</sub>)<sub>n</sub>-N(R<sup>4</sup>-(CH<sub>2</sub>)<sub>n</sub>-Z)(R<sup>5</sup>-(CH<sub>2</sub>)<sub>n</sub>-Z), (CH<sub>2</sub>)<sub>n</sub>-Z, O-(CH<sub>2</sub>)<sub>n</sub>-Z, (CH<sub>2</sub>)<sub>n</sub>-O-Z, S-(CH<sub>2</sub>)<sub>n</sub>-Z, CH=CH-Z, 1,2-methylenedioxy, C(O)OH, C(O)O-[(C<sub>1</sub>-C<sub>6</sub>)-straight or branched alkyl], C(O)O-(CH<sub>2</sub>)<sub>n</sub>-Z or C(O)-N(R<sup>4</sup>)(R<sup>5</sup>);

wherein each of R<sup>4</sup> and R<sup>5</sup> are independently  
15 hydrogen, (C<sub>1</sub>-C<sub>6</sub>)-straight or branched alkyl, (C<sub>3</sub>-C<sub>5</sub>)-straight or branched alkenyl, or wherein R<sup>4</sup> and R<sup>5</sup>, when bound to the same nitrogen atom, are taken together with the nitrogen atom to form a 5 or 6 membered ring, wherein said ring optionally contains 1 to 3 additional  
20 heteroatoms independently selected from N, O or S; wherein said alkyl, alkenyl or alkynyl groups in R<sub>4</sub> and R<sub>5</sub> are optionally substituted with Z.

each n is independently 0 to 4;

each Z is independently selected from a saturated,  
25 partially saturated or unsaturated, monocyclic or bicyclic ring system, wherein each ring comprises 5 to 7 ring atoms independently selected from C, N, O or S; and wherein no more than 4 ring atoms are selected from N, O or S;

30 wherein 1 to 4 hydrogen atoms in Z are optionally and independently replaced with halo, hydroxy, nitro, cyano, C(O)OH, (C<sub>1</sub>-C<sub>3</sub>)-straight or branched alkyl,

O-(C<sub>1</sub>-C<sub>3</sub>)-straight or branched alkyl,  
C(O)O-[(C<sub>1</sub>-C<sub>3</sub>)-straight or branched alkyl], amino,  
NH[(C<sub>1</sub>-C<sub>3</sub>)-straight or branched alkyl], or  
N-[(C<sub>1</sub>-C<sub>3</sub>)-straight or branched alkyl]<sub>2</sub>;

5 K is selected from E, (C<sub>1</sub>-C<sub>6</sub>)-straight or branched  
alkyl, (C<sub>2</sub>-C<sub>6</sub>)-straight or branched alkenyl or alkynyl,  
wherein 1 to 2 hydrogen atoms in said alkyl, alkenyl or  
alkynyl is optionally and independently replaced with E;  
wherein K is optionally substituted with up to 3  
10 substituents selected from halogen, OH, O-(C<sub>1</sub>-C<sub>6</sub>)-alkyl,  
O-(CH<sub>2</sub>)<sub>n</sub>-Z, NO<sub>2</sub>, CO<sub>2</sub>H, C(O)-O-(C<sub>1</sub>-C<sub>6</sub>)-alkyl, C(O)NR<sup>4</sup>R<sup>5</sup>,  
NR<sup>4</sup>R<sup>5</sup> and (CH<sub>2</sub>)<sub>n</sub>-Z; or,

G, when present, is -S(O)<sub>2</sub>-, -C(O)-, -S(O)<sub>2</sub>-Y-,  
-C(O)-Y-, -C(O)-C(O)-, or -C(O)-C(O)-Y-;

15 Y is oxygen, or N(R<sup>6</sup>);

wherein R<sup>6</sup> is hydrogen, E, (C<sub>1</sub>-C<sub>6</sub>)-straight or  
branched alkyl, (C<sub>3</sub>-C<sub>6</sub>)-straight or branched alkenyl or  
alkynyl; or wherein R<sup>6</sup> and D are taken together with the  
atoms to which they are bound to form a 5 to 7 membered  
20 ring system wherein said ring optionally contains 1 to 3  
additional heteroatoms independently selected from O, S,  
N, NH, SO, or SO<sub>2</sub>; and wherein said ring is optionally  
benzofused;

D is hydrogen, (C<sub>1</sub>-C<sub>7</sub>)-straight or branched  
25 alkyl, (C<sub>2</sub>-C<sub>7</sub>)-straight or branched alkenyl or alkynyl,  
(C<sub>5</sub>-C<sub>7</sub>)-cycloalkyl or cycloalkenyl optionally substituted  
with (C<sub>1</sub>-C<sub>6</sub>)-straight or branched alkyl or (C<sub>2</sub>-C<sub>7</sub>)-straight  
or branched alkenyl or alkynyl, [(C<sub>1</sub>-C<sub>7</sub>)-alkyl]-E,  
[(C<sub>2</sub>-C<sub>7</sub>)-alkenyl or alkynyl]-E, or E;

30 wherein 1 to 2 of the CH<sub>2</sub> groups of said alkyl,  
alkenyl or alkynyl chains in D is optionally replaced by  
-O-, -S-, -S(O)-, -S(O<sub>2</sub>)-, or -N(R<sup>3</sup>);

provided that when G is selected from  $-S(O)_2-$ ,  $-C(O)C(O)-$ ,  $-SO_2-Y$ , or  $-C(O)-Y$ , or  $-C(O)C(O)-Y$ , wherein  $Y = O$ ; then D is not hydrogen; and  $x = 0$  or 1.

5

2. The compound according to claim 1, wherein:

each of A and B is independently selected from  $-CH_2-CH_2-E$  or  $-CH_2-CH_2-CH_2-E$ ; and

10 E is a monocyclic or bicyclic aromatic ring system, wherein said ring comprises 5-7 ring atoms independently selected from C, N, O or S, and wherein 1 to 4 ring atoms are independently selected from N, O or S;

wherein 1 to 4 hydrogen atoms in E are optionally  
15 and independently replaced with halogen, hydroxyl, hydroxymethyl, nitro,  $SO_3H$ , trifluoromethyl, trifluoromethoxy,  $(C_1-C_6)$ -straight or branched alkyl,  $(C_2-C_6)$ -straight or branched alkenyl,  $O-[(C_1-C_6)$ -straight or branched alkyl],  $O-[(C_3-C_6)$ -straight or branched  
20 alkenyl],  $(CH_2)_n-N(R^4)(R^5)$ ,  $(CH_2)_n-NH(R^4)-(CH_2)_n-Z$ ,  $(CH_2)_n-N(R^4-(CH_2)_n-Z)(R^5-(CH_2)_n-Z)$ ,  $(CH_2)_n-Z$ ,  $O-(CH_2)_n-Z$ ,  $(CH_2)_n-O-Z$ ,  $S-(CH_2)_n-Z$ ,  $CH=CH-Z$ , 1,2-methylenedioxy,  $C(O)OH$ , or  $C(O)-N(R^4)(R^5)$ .

25 3. The compound according to claim 1 or 2, wherein D is an aromatic monocyclic or bicyclic ring system, wherein each ring comprises 5 to 7 ring atoms independently selected from C, N, O or S; and wherein no more than 4 ring atoms are selected from N, O or S.

30

4. The compound according to claim 3, wherein D is substituted phenyl.

5. The compound according to claim 4, wherein  
G is  $-C(O)C(O)-$ .

5 6. The compound according to claim 4, wherein  
G is  $-SO_2-$ .

7. The compound according to claim 4, wherein  
G is  $-C(O)-$ .

10 8. The compound according to claim 4, wherein  
G is  $-C(O)Y-$ .

9. A compound selected from compound nos. 1  
15 and 2.

10. The compound according to claim 2, wherein  
each of A and B is independently selected from  $-CH_2-CH_2-E$   
or  $-CH_2-CH_2-CH_2-E$ ; and  
20 E is pyridyl.

11. A composition comprising a compound  
according to claim 1 and a pharmaceutically effective  
carrier.

25 12. The composition according to claim 12,  
further comprising a neurotrophic factor.

13. The composition according to claim 13,  
30 wherein said neurotrophic factor is selected from nerve  
growth factor (NGF), insulin-like growth factor (IGF-1)  
and its active truncated derivatives such as gIGF-1 and

Des(1-3)IGF-I, acidic and basic fibroblast growth factor (aFGF and bFGF, respectively), platelet-derived growth factors (PDGF), brain-derived neurotrophic factor (BDNF), ciliary neurotrophic factors (CNTF), glial cell line-  
5 derived neurotrophic factor (GDNF), neurotrophin-3 (NT-3) and neurotrophin 4/5 (NT-4/5).

14 15. The composition according to claim 14,  
wherein said neurotrophic factor is nerve growth factor  
10 (NGF).

15 16. A method for stimulating neuronal  
regeneration in a patient or in an ex vivo nerve cell,  
comprising the step of administering to said patient or  
15 said nerve cell a compound according to any one of claims  
1-12.

16 17. The method according to claim 16, wherein  
said compound is administered to a patient and is  
20 formulated together with a pharmaceutically suitable  
carrier into a pharmaceutically acceptable composition.

17 18. The method according to claim 17,  
comprising the additional step of administering to said  
25 patient a neurotrophic factor either as part of a  
multiple dosage form together with said compound or as a  
separate dosage form.

18 19. The method according to claim 18, wherein  
30 said neurotrophic factor is selected from nerve growth  
factor (NGF), insulin-like growth factor (IGF-1) and its  
active truncated derivatives such as gIGF-1 and

Des(1-3)IGF-I, acidic and basic fibroblast growth factor (aFGF and bFGF, respectively), platelet-derived growth factors (PDGF), brain-derived neurotrophic factor (BDNF), ciliary neurotrophic factors (CNTF), glial cell line-  
5 derived neurotrophic factor (GDNF), neurotrophin-3 (NT-3) and neurotrophin 4/5 (NT-4/5).

19 20. The method according to claim 19, wherein said neurotrophic factor is nerve growth factor (NGF).

10 20 21. The method according to claim 16, wherein said method is used to treat a patient suffering from a disease selected from trigeminal neuralgia, glossopharyngeal neuralgia, Bell's Palsy, myasthenia  
15 gravis, muscular dystrophy, muscle injury, progressive muscular atrophy, progressive bulbar inherited muscular atrophy, herniated, ruptured, or prolapsed intervertebrae disk syndrome's, cervical spondylosis, plexus disorders, thoracic outlet destruction syndromes, peripheral  
20 neuropathies, such as those caused by lead, dapsone, ticks, or porphyria, other peripheral myelin disorders, Alzheimer's disease, Gullain-Barre syndrome, Parkinson's disease and other Parkinsonian disorders, ALS, Tourette's syndrome, multiple sclerosis, other central myelin  
25 disorders, stroke and ischemia associated with stroke, neural paropathy, other neural degenerative diseases, motor neuron diseases, sciatic crush, neuropathy associated with diabetes, spinal cord injuries, facial nerve crush and other trauma, chemotherapy- and other  
30 medication-induced neuropathies, and Huntington's disease.

21 22. The method according to claim 16,  
wherein said method is used to stimulate neuronal  
regeneration in an ex vivo nerve cell.

5 22 23. The method according to claim 22,  
comprising the additional step of contacting said ex  
vivo nerve cell with a neurotrophic factor.

23 24. The method according to claim 23, wherein  
10 said neurotrophic factor is selected from nerve growth  
factor (NGF), insulin-like growth factor (IGF-1) and its  
active truncated derivatives such as gIGF-1 and  
Des(1-3)IGF-I, acidic and basic fibroblast growth factor  
(aFGF and bFGF, respectively), platelet-derived growth  
15 factors (PDGF), brain-derived neurotrophic factor (BDNF),  
ciliary neurotrophic factors (CNTF), glial cell line-  
derived neurotrophic factor (GDNF), neurotrophin-3 (NT-  
3) and neurotrophin 4/5 (NT-4/5).

20 24 25. The method according to claim 24, wherein  
said neurotrophic factor is nerve growth factor (NGF).